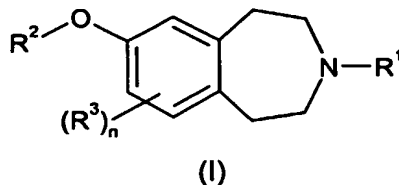


CLAIMS:

1. A compound of formula (I) or a pharmaceutically acceptable salt thereof:



wherein:

R¹ represents -C₃₋₇ cycloalkyl optionally substituted by C₁₋₃ alkyl;

R² represents hydrogen, -C₁₋₆ alkyl, -X-C₃₋₈ cycloalkyl, -X-aryl, -X-heterocyclyl, -X-heteroaryl, -X-C₃₋₈ cycloalkyl-Y-C₃₋₈ cycloalkyl, -X-C₃₋₈ cycloalkyl-Y-aryl, -X-C₃₋₈ cycloalkyl-Y-heteroaryl, -X-C₃₋₈ cycloalkyl-Y-heterocyclyl, -X-aryl-Y-C₃₋₈ cycloalkyl, -X-aryl-Y-aryl, -X-aryl-Y-heteroaryl, -X-aryl-Y-heterocyclyl, -X-heteroaryl-Y-C₃₋₈ cycloalkyl, -X-heteroaryl-Y-aryl, -X-heteroaryl-Y-heteroaryl, -X-heteroaryl-Y-heterocyclyl, -X-heterocyclyl-Y-C₃₋₈ cycloalkyl, -X-heterocyclyl-Y-aryl, -X-heterocyclyl-Y-heteroaryl, -X-heterocyclyl-Y-heterocyclyl;

X represents a bond or C₁₋₆ alkyl;

Y represents a bond, C₁₋₆ alkyl, CO, COC₂₋₆ alkenyl, O or SO₂;

R³ represents halogen, C₁₋₆ alkyl, C₁₋₆ alkoxy, cyano, amino or trifluoromethyl;

n is 0, 1 or 2;

wherein said alkyl, cycloalkyl, aryl, heteroaryl and heterocyclyl groups of R² may be optionally substituted by one or more substituents (eg. 1, 2 or 3) which may be the same or different, and which are selected from the group consisting of halogen, hydroxy, cyano, nitro, =O, trifluoromethyl, trifluoromethoxy, fluoromethoxy, difluoromethoxy, C₁₋₆ alkyl, pentafluoroethyl, C₁₋₆ alkoxy, arylC₁₋₆ alkoxy, C₁₋₆ alkylthio, C₁₋₆ alkoxyC₁₋₆ alkyl, C₃₋₇ cycloalkylC₁₋₆ alkoxy, C₁₋₆ alkanoyl, C₁₋₆ alkoxy carbonyl, C₁₋₆ alkylsulfonyl, C₁₋₆ alkylsulfinyl, C₁₋₆ alkylsulfonyloxy, C₁₋₆ alkylsulfonylC₁₋₆ alkyl, sulfonyl, arylsulfonyl, arylsulfonyloxy, arylsulfonylC₁₋₆ alkyl, aryloxy, C₁₋₆ alkylsulfonamido, C₁₋₆ alkylamino, C₁₋₆ alkylamido, -R⁴, -CO₂R⁴, -COR⁴, C₁₋₆ alkylsulfonamidoC₁₋₆ alkyl, C₁₋₆ alkylamidoC₁₋₆ alkyl, arylsulfonamido, arylcarboxamido, arylsulfonamidoC₁₋₆ alkyl, arylcarboxamidoC₁₋₆ alkyl, aroyl, aroylC₁₋₆ alkyl, arylC₁₋₆ alkanoyl, or a group -NR⁵R⁶, -C₁₋₆ alkyl-NR⁵R⁶, -C₃₋₈ cycloalkyl-NR⁵R⁶, -CONR⁵R⁶, -NR⁵COR⁶, -NR⁵SO₂R⁶, -OCONR⁵R⁶, -NR⁵CO₂R⁶, -NR⁴CONR⁵R⁶ or -SO₂NR⁵R⁶ (wherein R⁴, R⁵ and R⁶ independently represent hydrogen, C₁₋₆ alkyl, -C₃₋₈ cycloalkyl, -C₁₋₆ alkyl-C₃₋₈ cycloalkyl, aryl, heterocyclyl or heteroaryl or -NR⁵R⁶ may represent a nitrogen containing heterocyclyl group, wherein said R⁴, R⁵ and R⁶ groups may be optionally substituted by one or more substituents (eg. 1, 2 or 3) which may be the same or different, and which are selected from the group consisting of halogen, hydroxy, C₁₋₆ alkyl, C₁₋₆ alkoxy, cyano, amino, =O or trifluoromethyl); or solvates thereof.

2. A compound of formula (I) as defined in claim 1 wherein R¹ represents unsubstituted cyclobutyl, cyclopentyl or cyclohexyl.

3. A compound of formula (I) as defined in claim 2 wherein R¹ represents unsubstituted cyclobutyl.

4. A compound of formula (I) as defined in any one of claims 1 to 3 wherein R² represents:

hydrogen;

-C₁₋₆ alkyl optionally substituted by a -CO₂R⁴ or -CONR⁵R⁶ group;

-X-C₃₋₈ cycloalkyl-Y-heterocyclyl;

-X-aryl-Y-C₃₋₈ cycloalkyl;

-X-aryl optionally substituted by one or two halogen, C₁₋₆ alkyl, C₁₋₆ alkoxy, -CO₂R⁴, -CONR⁵R⁶, -NR⁵COR⁶, -SO₂NR⁵R⁶ or cyano groups;

-X-aryl-Y-heterocyclyl optionally substituted by one or two =O, halogen or R⁴ groups;

-X-heteroaryl optionally substituted by one or two halogen, C₁₋₆ alkyl, C₁₋₆ alkoxy, cyano, nitro, -OR⁴, -COR⁴, -CO₂R⁴, -NR⁵R⁶, -NR⁵COR⁶, -CONR⁵R⁶ or =O groups;

-X-heteroaryl-Y-aryl optionally substituted by a C₁₋₆ alkylsulfonyl or -NR⁵COR⁶

group;

-X-heteroaryl-Y-heteroaryl optionally substituted by a C₁₋₆ alkyl group;

-X-heteroaryl-Y-heterocyclyl optionally substituted by one or two =O, C₁₋₆ alkyl, -OR⁴ or halogen groups;

-X-heteroaryl-Y-heterocyclyl optionally substituted by one or two =O, C₁₋₆ alkyl, -OR⁴

or halogen groups;

-X-heterocyclyl optionally substituted by a C₁₋₆ alkylsulfonyl, C₁₋₆ alkoxycarbonyl, -CO₂R⁴, -COR⁴ or -COR⁵R⁶ group;

-X-heterocyclyl-Y-aryl optionally substituted by a halogen, cyano, C₁₋₆ alkylsulfonyl, R⁴ or -CONR⁵R⁶ group;

-X-heterocyclyl-Y-heterocyclyl optionally substituted by one or two =O or R⁴ groups;

-X-heterocyclyl-Y-C₃₋₈ cycloalkyl; or

-X-heterocyclyl-Y-heteroaryl optionally substituted by one or two C₁₋₆ alkyl, =O, cyano or -CONR⁵R⁶ groups.

5. A compound of formula (I) as defined in claim 4 wherein R² represents

-X-aryl optionally substituted by one or two halogen, C₁₋₆ alkyl, C₁₋₆ alkoxy, -CO₂R⁴, -CONR⁵R⁶, -NR⁵COR⁶, -SO₂NR⁵R⁶ or cyano groups;

-X-aryl-Y-heterocyclyl optionally substituted by one or two =O, halogen or R⁴ groups;

-X-heteroaryl optionally substituted by one or two halogen, C₁₋₆ alkyl, C₁₋₆ alkoxy, cyano, nitro, -OR⁴, -CO₂R⁴, -COR⁴, -NR⁵R⁶, -NR⁵COR⁶, -CONR⁵R⁶ or =O groups;

-X-heteroaryl-Y-heterocyclyl optionally substituted by one or two =O, C₁₋₆ alkyl, -OR⁴ or halogen groups; or

-X-heterocyclyl-Y-heterocyclyl optionally substituted by one or two =O or R⁴ groups.

5 6. A compound of formula (I) as defined in claim 5 wherein R² represents:

-X-aryl optionally substituted by one or two halogen, C₁₋₆ alkoxy, -CONR⁵R⁶, -NR⁵COR⁶ or cyano groups;

-X-aryl-Y-heterocyclyl optionally substituted by one or two =O or halogen groups; unsubstituted -X-heterocyclyl-Y-heterocyclyl;

10 -X-heteroaryl optionally substituted by CON(H)(Me)); or

-X-heteroaryl-Y-heterocyclyl wherein said heterocyclyl group is optionally substituted by an =O group.

15 7. A compound of formula (I) as defined in claim 6 wherein R² represents:

-phenyl optionally substituted by one or two fluorine, methoxy, -CON(H)(Me), -NHCOMe or cyano groups;

-phenyl-pyrrolidinyl optionally substituted by one or two =O or fluorine groups; unsubstituted -piperidinyl-CO-morpholinyl;

-2-pyridinyl or -2-pyrazinyl optionally substituted by CON(H)(Me); or

20 -2-pyridinyl-N-pyrrolidinyl wherein said pyrrolidinyl group is optionally substituted by an =O group.

25 8. A compound of formula (I) as defined in claim 7 wherein R² represents 4-methylaminocarbonylpyridin-2-yl.

9. A compound of formula (I) as defined in any one of claims 1 to 6 wherein X represents a bond or -CH₂-.

30 10. A compound of formula (I) as defined in claim 9 wherein X represents a bond.

11. A compound of formula (I) as defined in any one of claims 1 to 6, 9 or 10 wherein Y represents a bond, CO, SO₂ or -CO-CH=CH-.

35 12. A compound of formula (I) as defined in claim 11 wherein Y represents a bond.

13. A compound of formula (I) as defined in any one of claims 1 to 5 or 9 to 12 wherein R⁴ represents hydrogen, C₁₋₆ alkyl, -C₁₋₆ alkyl-C₃₋₈ cycloalkyl, aryl, heterocyclyl or heteroaryl optionally substituted by a halogen or C₁₋₆ alkoxy group.

40 14. A compound of formula (I) as defined in any one of claims 1 to 5 or 9 to 12 wherein R⁵ and R⁶ independently represent hydrogen, C₁₋₆ alkyl, -C₃₋₈ cycloalkyl, -C₁₋₆ alkyl-C₃₋₈ cycloalkyl, aryl or heterocyclyl optionally substituted by a halogen, cyano or C₁₋₆ alkoxy

group or $\text{--NR}^5\text{R}^6$ represents a nitrogen containing heterocyclyl group optionally substituted by one or two =O groups.

15. A compound of formula (I) as defined in claim 14 wherein R^5 and R^6 independently represent hydrogen, C_{1-6} alkyl, --C_{3-8} cycloalkyl or --C_{1-6} alkyl- C_{3-8} cycloalkyl.

16. A compound of formula (I) as defined in any one of claims 1 to 15 wherein n represents 0 or 1.

17. A compound of formula (I) as defined in claim 16 wherein n represents 1 and R^3 represents a halogen atom or a cyano group.

18. A compound of formula (I) as defined in claim 16 wherein n represents 0.

19. A compound according to claim 1 which is a compound of formula E1-E288 or a pharmaceutically acceptable salt thereof.

20. A compound of formula (I) as defined in claim 1 which is:
5-(3-Cyclobutyl-2,3,4,5-tetrahydro-1*H*-benzo[d]azepin-7-yloxy)-pyrazine-2-carboxylic acid methyl amide; or
1-{6-[(3-Cyclobutyl-2,3,4,5-tetrahydro-1*H*-3-benzazepin-7-yl)oxy]-3-pyridinyl}-2-pyrrolidinone;
or a pharmaceutically acceptable salt thereof.

21. A compound of formula (I) as defined in claim 1 which is 6-(3-cyclobutyl-2,3,4,5-tetrahydro-1*H*-benzo[d]azepin-7-yloxy)-*N*-methyl-nicotinamide or a pharmaceutically acceptable salt thereof.

22. A pharmaceutical composition which comprises the compound of formula (I) as defined in any one of claims 1 to 21 or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable carrier or excipient.

23. A compound as defined in any one of claims 1 to 21 for use in therapy.

24. A compound as defined in any one of claims 1 to 21 for use in the treatment of neurological diseases.

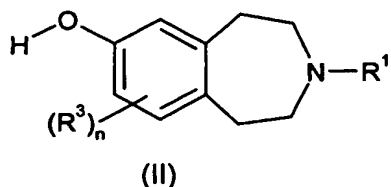
25. Use of a compound as defined in any one of claims 1 to 21 in the manufacture of a medicament for the treatment of neurological diseases.

26. A method of treatment of neurological diseases which comprises administering to a host in need thereof an effective amount of a compound of formula (I) as defined in any one of claims 1 to 21 or a pharmaceutically acceptable salt thereof.

27. A pharmaceutical composition for use in the treatment of neurological diseases which comprises the compound of formula (I) as defined in any one of claims 1 to 21 or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable carrier.

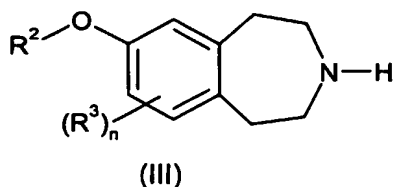
28. A process for the preparation of a compound of formula (I) or a pharmaceutically acceptable salt thereof, which process comprises:

(a) reacting a compound of formula (II)



wherein R^1 , R^3 and n are as defined in claim 1, with a compound of formula R^2-L^1 , wherein R^2 is as defined in claim 1 for R^2 or a group convertible thereto and L^1 represents a suitable leaving group such as a halogen atom or an optionally activated hydroxyl group;

(b) reacting a compound of formula (III)



wherein R^2 , R^3 and n are as defined in claim 1, with a compound of formula $R^{1'}-L^2$, wherein $R^{1'}$ is as defined in claim 1 for R^1 or a group convertible thereto and L^2 represents a suitable leaving group such as a halogen atom; or

(c) reacting a compound of formula (III) as defined in claim 1, with a ketone of formula $R^{1'}=O$, wherein $R^{1'}$ is as defined in claim 1 for R^1 or a group convertible thereto; or

(d) deprotecting a compound of formula (I) which is protected; and

(e) interconversion to other compounds of formula (I).